

PRIOR-AUTHORIZATION OF LENALIDOMIDE (REVLIMID™)

Maryland Pharmacy Program

Tel#: 410-767-1455 or 1-800-492-5231 Option 3-Fax form to: 410-333-5398

(Incomplete forms will be returned)

Patient Information

Patient location: ____ Home; ____ Hospital ____ Clinic ____ Office Age: ____ Date of Birth: ____/____/____
Patient Name: _____
MA ID#: _____ Address: _____
Tel.#:(____)____-_____
Has Patient received Red Blood Cell (RBC) transfusions in the past? ☐ Yes ☐ No
Is Patient currently receiving RBC transfusions? ☐ Yes ☐ No
Is Prescriber registered in the Rev Assist program? ☐ Yes ☐ No
What are the prior antimyeloma therapies tried? _____

Attach a dated copy of the most recent lab tests and provide test results for the following:

Platelet count: _____/mcL Absolute neutrophil count (ANC) _____/mcL
Serum creatinine: _____mg/dL; SGOT/AST or SGPT/ALT _____ Direct bilirubin: _____mg/dL

Prescriber Information

Is Revlimid™ prescribed as part of a clinical study? ☐ Yes ☐ No
Specify sponsoring organization/drug manufacturer _____
Specify purpose of study: _____

Note: Off-label use or use of Revlimid™ at dosages other than recommended by FDA must be medically necessary and supported by/ documented in one of the three official compendia, the American Hospital Formulary Service Drug Information, the Micromedex/Drugdex drug database, and the U.S. Pharmacopeia.

I certify that Patient is not enrolled in any study involving the requested drug. I will be supervising the patient's treatment accordingly. Supporting medical documentation is kept on file in the patient's medical record.

_____, M.D. Prescriber's Name: _____ Date: _____
(Prescriber's signature) Tel# (____) - ____ - ____ Fax# (____) - ____ - ____
License #: _____ DEA #: _____ Specialty : _____
Address: _____

Prescription Information

Drug/Strength/dosage prescribed: _____
List diagnosis for which the drug was prescribed:
☐ Treatment of transfusion-dependent anemia due to Low-or Intermediate-1 risk myelodysplastic syndromes (MDS) associated with a deletion 5q cytogenetic abnormality with or without transient tyrosinemia of the newborn (TTN)
☐ MDS not related to 5 q deletion abnormality
☐ Multiple myeloma
☐ Other: _____

FOR INTERNAL USE

Approved: ☐ Denied: ☐ Date: _____ Reviewer's Initials _____
Reason for denial: _____
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